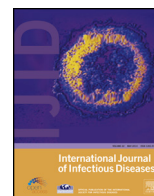




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Medical Imagery

Oculomotor nerve palsy in invasive intracranial aspergillosis

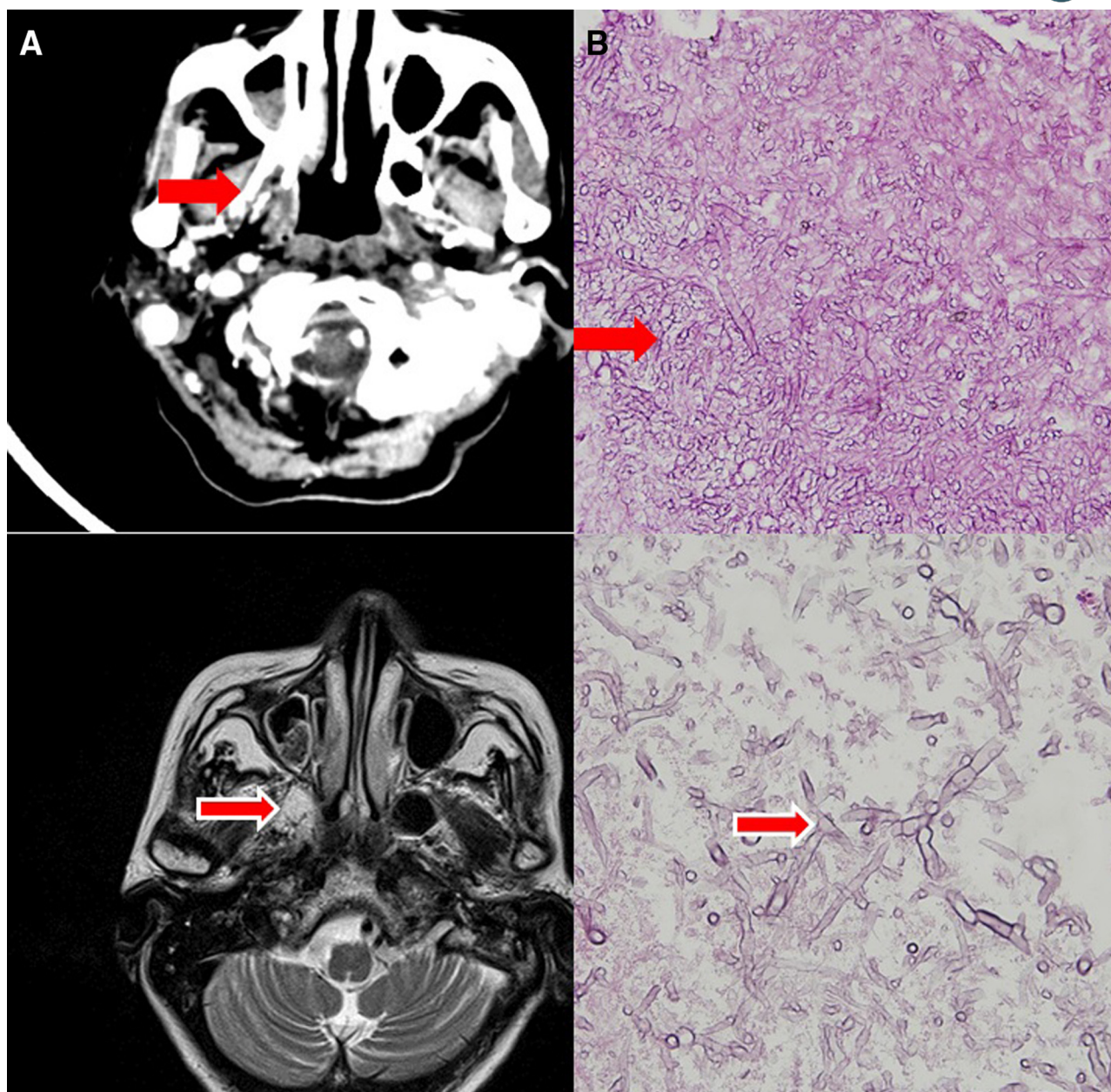


Figure 1. A Contrast-enhanced computed tomography scan of the brain demonstrating abnormal contrast enhancement of the maxillary sinus, cavernous sinus, and superior orbital apex on the axial image (red arrow). Axial T2-weighted contrast-enhanced magnetic resonance imaging of the brain showing abnormal contrast enhancement of the maxillary sinus, cavernous sinus, and superior orbital apex, with skull base bony destruction (red–white arrow). (B) Photomicrograph revealing a fungal ball with numerous branching hyphae (hematoxylin–eosin stain, $\times 100$, red arrow) and characteristic septate fungal hyphae of an *Aspergillus* species, and hyphae branching dichotomously at a 45-degree angle (hematoxylin–eosin stain, $\times 400$, red–white arrow).

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The patient was a 70-year-old diabetic Taiwanese woman, with a past medical history of otitis media status post surgery performed 2 years prior. She presented to the emergency department with a progression of headache, double vision, and ptosis of the right eye occurring over the course of 1 month, following an inability to perform lateral gaze. A neurological examination demonstrated palsy of intracranial nerves III, IV, and VI of the right eye. A computed tomography scan and magnetic resonance imaging of the brain showed abnormal contrast enhancement of the right maxillary sinus, cavernous sinus, and superior orbital apex, along with skull base bony destruction (Figure 1A). A biopsy of the right paranasal sinus was performed by endoscopic sinus surgery, and photomicrographs revealed a fungal ball with numerous characteristic septate fungal hyphae of an *Aspergillus* species and hyphae branched dichotomously at a 45-degree angle (Figure 1B). Treatment with intravenous amphotericin B (50 mg daily) was prescribed for 1 week; this was then changed to voriconazole (200 mg twice daily) for 6 months, in the outpatient department.

Invasive aspergillosis has been shown to occur in 10.7% of patients with acute myeloid leukemia receiving induction chemotherapy.¹ Invasive aspergillosis of the intracranial nerve system is a difficult diagnostic challenge and has a high mortality rate, especially in those in an immunocompromised condition. Sinusitis, vascular narrowing or obstruction, and bony destruction are radiological signs of invasive fungal infection. A definite diagnosis of aspergillosis is based on the histopathological characteristics of the tissue involved demonstrating a fungal ball with aggregates of spores and hyphae with 45-degree branching.^{2–4} Surgical resection of the focal lesion remains the cornerstone of management, followed by systemic antifungal medications.^{4,5}

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Consent: Written informed consent was obtained from the patient for publication of this case report and all accompanying images.

Conflict of interest: No competing interest declared.

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